

WEST Search History

DATE: Friday, September 21, 2007

Hide?	Set Name	Query	Hit Count
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DB=USPT; PLUR=YES; OP=OR

<input type="checkbox"/>	L3	l1 and ergoline	0
<input type="checkbox"/>	L2	L1 and cabergoline	0
<input type="checkbox"/>	L1	546/75	52

END OF SEARCH HISTORY

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LOGINID:SSSPTA1612RXD

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

* * * * * Welcome to STN International * * * * *

NEWS	1		Web Page for STN Seminar Schedule - N. America
NEWS	2	JUL 02	LMEDLINE coverage updated
NEWS	3	JUL 02	SCISEARCH enhanced with complete author names
NEWS	4	JUL 02	CHEMCATS accession numbers revised
NEWS	5	JUL 02	CA/Capplus enhanced with utility model patents from China
NEWS	6	JUL 16	Capplus enhanced with French and German abstracts
NEWS	7	JUL 18	CA/Capplus patent coverage enhanced
NEWS	8	JUL 26	USPATFULL/USPAT2 enhanced with IPC reclassification
NEWS	9	JUL 30	USGENE now available on STN
NEWS	10	AUG 06	CAS REGISTRY enhanced with new experimental property tags
NEWS	11	AUG 06	BEILSTEIN updated with new compounds
NEWS	12	AUG 06	FSTA enhanced with new thesaurus edition
NEWS	13	AUG 13	CA/Capplus enhanced with additional kind codes for granted patents
NEWS	14	AUG 20	CA/Capplus enhanced with CAS indexing in pre-1907 records
NEWS	15	AUG 27	Full-text patent databases enhanced with predefined patent family display formats from INPADOCDB
NEWS	16	AUG 27	USPATOLD now available on STN
NEWS	17	AUG 28	CAS REGISTRY enhanced with additional experimental spectral property data
NEWS	18	SEP 07	STN AnaVist, Version 2.0, now available with Derwent World Patents Index
NEWS	19	SEP 13	FORIS renamed to SOFIS
NEWS	20	SEP 13	INPADOCDB enhanced with monthly SDI frequency
NEWS	21	SEP 17	CA/Capplus enhanced with printed CA page images from 1967-1998
NEWS	22	SEP 17	Capplus coverage extended to include traditional medicine patents
NEWS EXPRESS	19	SEPTEMBER 2007:	CURRENT WINDOWS VERSION IS V8.2, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 19 SEPTEMBER 2007.
NEWS HOURS	STN Operating Hours Plus Help Desk Availability		
NEWS LOGIN	Welcome Banner and News Items		
NEWS IPC8	For general information regarding STN implementation of IPC 8		

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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 14:13:59 ON 21 SEP 2007

=> file registry

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

0.42

0.42

FILE 'REGISTRY' ENTERED AT 14:14:59 ON 21 SEP 2007

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STRUCTURE FILE UPDATES: 20 SEP 2007 HIGHEST RN 947666-94-6

DICTIONARY FILE UPDATES: 20 SEP 2007 HIGHEST RN 947666-94-6

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH June 29, 2007

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=> s cageline/cn

L1 0 CAGELINE/CN

=> s cabergoline/cn

L2 1 CABERGOLINE/CN

=> d

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2007 ACS on STN

RN 81409-90-7 REGISTRY

ED Entered STN: 16 Nov 1984

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-[(ethylamino)carbonyl]-6-(2-propenyl)-, (8 β)- (9CI)

OTHER NAMES:

CN Cabaser

CN Cabergoline

CN Dostinex

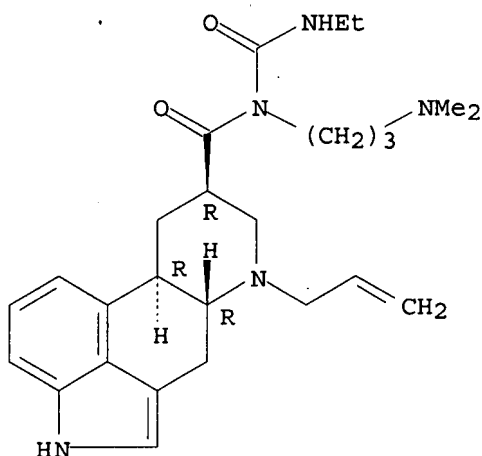
CN Galastop

CN Sogilen

FS STEREOSEARCH
 MF C26 H37 N5 O2
 CI COM

LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CBNB, CHEMCATS, CIN, CSCHEM, DDFU, DRUGU, EMBASE, IMSCSEARCH, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*, PATDPASPC, PHAR, PROMT, PROUSDDR, PS, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL, VETU
 (*File contains numerically searchable property data)
 Other Sources: WHO

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

418 REFERENCES IN FILE CA (1907 TO DATE)
 7 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
 419 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
12.30	12.72

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 14:15:32 ON 21 SEP 2007

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FILE COVERS 1907 - 21 Sep 2007 VOL 147 ISS 14
FILE LAST UPDATED: 20 Sep 2007 (20070920/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/infopolicy.html>

=> s 12

L3 419 L2

=> s 13 and heptane

62069 HEPTANE

1222 HEPTANES

62692 HEPTANE

(HEPTANE OR HEPTANES)

L4 8 L3 AND HEPTANE

=> d abs hitstr fbib 1-8

L4 ANSWER 1 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

AB A method of preparing cabergoline Form I, comprising forming a solvate including cabergoline and a p-disubstituted benzene of formula (A) or 1,3,5-trimethylbenzene and obtaining cabergoline Form I from the solvate. Another aspect of the present invention provides a method for preparing cabergoline Form I comprising dissolving cabergoline in p-disubstituted benzene or 1,3,5-trimethylbenzene and recovering the cabergoline Form I polymorph, suitably by direct crystallization of Form I or by recovery of a solvate which can be converted to Form I. Another aspect of the present invention provides a novel cabergoline polymorph designated cabergoline Form FB and a method of preparing said polymorph by dissolving or forming a solvate of cabergoline in fluorobenzene and recovering cabergoline Form FB.

IT 81409-90-7, Cabergoline

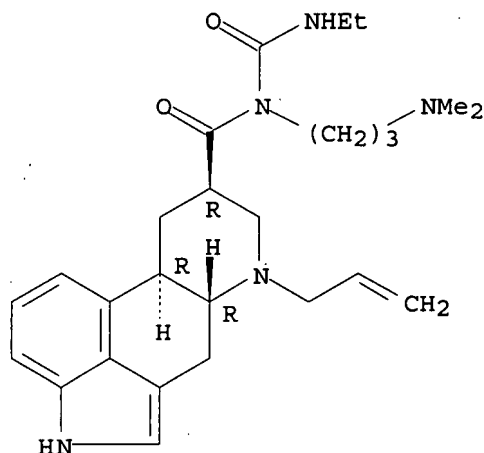
RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); RCT (Reactant); THU (Therapeutic use); BIOL (Biological study); PROC (Process); RACT (Reactant or reagent); USES (Uses)

(preparation of cabergoline in different crystal forms from various solvents)

RN 81409-90-7 CAPLUS

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2006:1012463 CAPLUS
 DN 145:383503
 TI Preparation of cabergoline in different crystal forms from various solvents
 IN Greenwood, Alan Kenneth; Mchattie, Derek; Bhatarah, Parveen; Aloui, Mahmoud
 PA UK
 SO U.S. Pat. Appl. Publ., 23pp., Cont.-in-part of U.S. Ser. No. 100,934.
 CODEN: USXXCO
 DT Patent
 LA English
 FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2006217408	A1	20060928	US 2005-268118	20051107
			GB 2005-5965	A 20050323
			US 2005-100934	A2 20050407
			GB 2005-15430	A 20050727
US 2006217555	A1	20060928	US 2005-100934	20050407
US 7238810	B2	20070703		
			GB 2005-5965	A 20050323
WO 2007012846	A1	20070201	WO 2006-GB2784	20060727
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			GB 2005-15430	A 20050727

PATENT FAMILY INFORMATION:

FAN 2006:1012462

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006100492	A2	20060928	WO 2006-GB1068	20060323
WO 2006100492	A3	20070104		

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GB 2005-5965 A 20050323

GB 2005-15430 A 20050727

WO 2007012846 A1 20070201 WO 2006-GB2784 20060727

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RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

GB 2005-15430 A 20050727

L4 ANSWER 2 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

AB A method of preparing cabergoline Form I, comprising forming a solvate including cabergoline and a p-disubstituted benzene or 1,3,5-trimethylbenzene and obtaining cabergoline Form I from the solvate. Another aspect of the present invention provides a method for preparing cabergoline Form I comprising dissolving cabergoline in p-disubstituted benzene or 1,3,5-trimethylbenzene and recovering the cabergoline Form I polymorph, suitably by direct crystallization of Form I or by recovery of a solvate which can be converted to Form I. Another aspect of the present invention provides a novel cabergoline polymorph designated cabergoline Form FB and a method of preparing said polymorph by dissolving or forming a solvate of cabergoline in fluorobenzene and recovering cabergoline Form FB.

IT 81409-90-7, Cabergoline

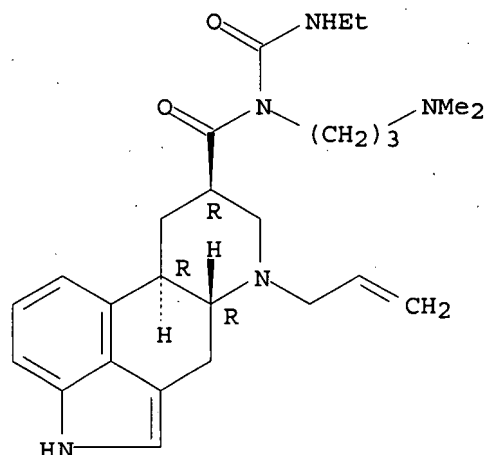
RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); RCT (Reactant); THU (Therapeutic use); BIOL (Biological study); PROC (Process); RACT (Reactant or reagent); USES (Uses)

(preparation of cabergoline in different crystal forms from various solvents)

RN 81409-90-7 CAPLUS

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2006:1012462 CAPLUS
DN 145:383502
TI Preparation of cabergoline in different crystal forms from various solvents
IN Greenwood, Alan, Kenneth; Mchattie, Derek; Bhatarah, Parveen; Aloui, Mahmoud
PA Resolution Chemicals Limited, UK
SO PCT Int. Appl., 44pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT. 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006100492	A2	20060928	WO 2006-GB1068	20060323
	WO 2006100492	A3	20070104		
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				GB 2005-5965	A 20050323
				GB 2005-15430	A 20050727
	WO 2007012846	A1	20070201	WO 2006-GB2784	20060727
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KG, KZ, MD, RU, TJ, TM

GB 2005-15430

A 20050727

PATENT FAMILY INFORMATION:

FAN 2006:1012463

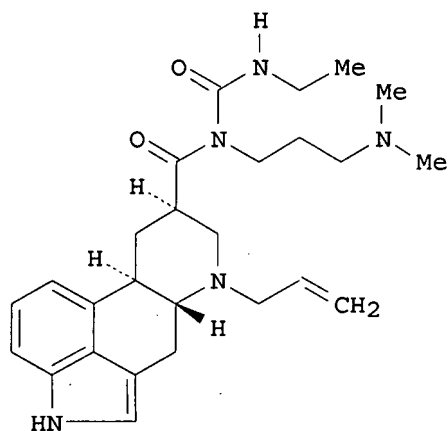
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PI	US 2006217408	A1	20060928	US 2005-268118	20051107
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				US 2005-100934	A2 20050407
				GB 2005-15430	A 20050727
	US 2006217555	A1	20060928	US 2005-100934	20050407
	US 7238810	B2	20070703		
				GB 2005-5965	A 20050323
	WO 2007012846	A1	20070201	WO 2006-GB2784	20060727
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GB 2005-15430

A 20050727

L4 ANSWER 3 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

GI



AB The present invention discloses a method for preparing cabergoline form 1 I by combining cabergoline and a solvent comprising ethylbenzene to form a solvate and obtaining form 1 from the solvate. Also disclosed is a method for preparing cabergoline form 1 by combining cabergoline and a first solvent to form a solution and addnl. including a second solvent to the solution, followed by crystallization to form cabergoline form 1. Further disclosed is a

solvate form of cabergoline comprising cabergoline and ethylbenzene and, optionally, n-heptane.

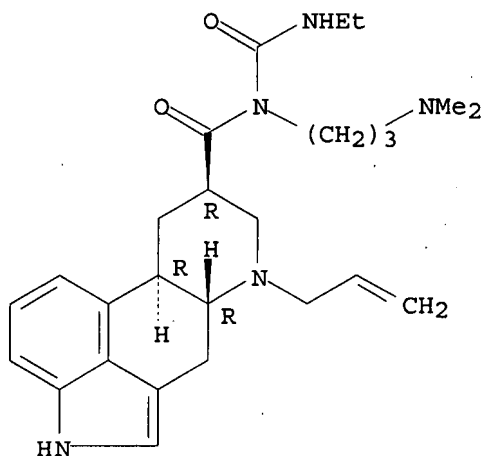
IT 81409-90-7P, Cabergoline

RL: PUR (Purification or recovery); PREP (Preparation)
(preparation of cabergoline from solvates)

RN 81409-90-7 CAPLUS

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-
[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2005:1168808 CAPLUS

DN 143:422511

TI Preparation of cabergoline and its solvate

IN Greenwood, Alan; Mchattie, Derek; Bhatarah, Parveen; Gamble, Mark

PA Resolution Chemicals Limited, UK

SO Eur. Pat. Appl., 9 pp.

CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1591445	A1	20051102	EP 2005-252709	20050429
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	US 2005245560	A1	20051103	US 2005-60991	20050218
	US 7186837	B2	20070306		
	AU 2005238276	A1	20051110	GB 2004-9785	A 20040430
				AU 2005-238276	20050429
				GB 2004-9785	A 20040430
				WO 2005-GB1649	W 20050429
	CA 2565440	A1	20051110	CA 2005-2565440	20050429
				GB 2004-9785	A 20040430
				WO 2005-GB1649	W 20050429
	WO 2005105796	A1	20051110	WO 2005-GB1649	20050429
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ,				

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 NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL,
 SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA,
 ZM, ZW
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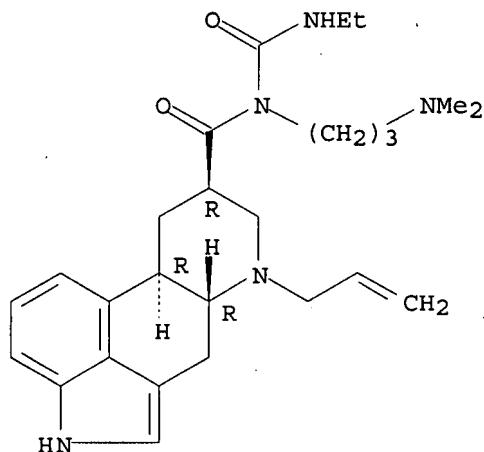
GB 2004-9785

A 20040430

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN
 AB New crystalline forms of cabergoline are disclosed. Also provided are novel
 processes for the preparation of cabergoline forms I, II, and VII, and
 amorphous cabergoline. Thus, cabergoline was dissolved in tert.-Bu Me
 ether and refluxed to give cabergoline tert.-Bu Me ether solvate form
 VIII.
 IT 81409-90-7, Cabergoline
 RL: FMU (Formation, unclassified); PRP (Properties); THU (Therapeutic
 use); BIOL (Biological study); FORM (Formation, nonpreparative); USES
 (Uses)
 (polymorphs of cabergoline)
 RN 81409-90-7 CAPLUS
 CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-
 [(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2004:1016006 CAPLUS
 DN 141:428033
 TI Polymorphs of cabergoline
 IN Cvak, Ladislav; Bednar, Roman; Sobotik, Roman; Jegorov, Alexandr
 PA Ivax Corporation, USA; Ivax Pharmaceuticals S.R.O
 SO PCT Int. Appl., 61 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

PATENT NO.

KIND

DATE

APPLICATION NO.

DATE

PI	WO 2004101510	A2	20041125	WO 2004-US14367	20040507
	WO 2004101510	A3	20050512		
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				US 2003-468887P	P 20030508
				US 2004-539494P	P 20040127
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	US 2005085499	A1	20050421	US 2004-841813	20040507
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				US 2004-539494P	P 20040127
				WO 2004-US14367	W 20040507
	JP 2007501274	T	20070125	JP 2006-532868	20040507
				US 2003-468887P	P 20030508
				US 2004-539494P	P 20040127
				WO 2004-US14367	W 20040507
	MX 2005PA11933	A	20060731	MX 2005-PA11933	20051107
				US 2003-468887P	P 20030508
				US 2004-539494P	P 20040127
				WO 2004-US14367	W 20040507

L4 ANSWER 5 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

AB The invention provides methods for preparing amorphous phys. form of cabergoline, and solvate form A of cabergoline useful in the preparation of the first mentioned phys. form. A method for treating a prolactin disorder with medicaments prepared from amorphous phys. form of cabergoline and solvate form A of cabergoline is also disclosed.

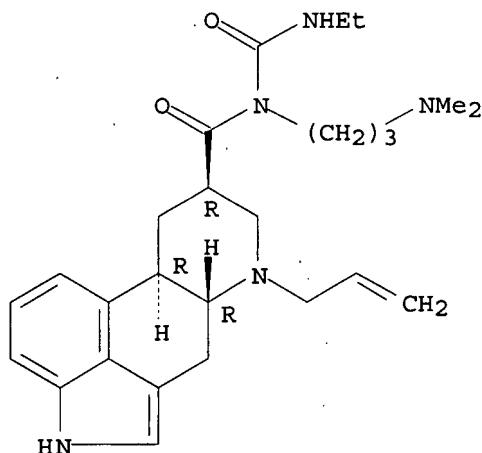
IT 81409-90-7P, Cabergoline
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (amorphous; tablets containing cabergoline or its solvate form for treatment of prolactin disorders)

RN 81409-90-7 CAPLUS

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-

[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2004:878162 CAPLUS
 DN 141:355382
 TI Preparation of cabergoline and its solvate form for treatment of prolactin disorders
 IN Gutman, Arie; Tishin, Boris; Vilenski, Alex; Agazade, Albay; Pertzikov, Boris; Nisnevich, Gennady
 PA Finetech Laboratories, Ltd., Israel
 SO U.S. Pat. Appl. Publ., 20 pp.
 CODEN: USXXCO
 DT Patent
 LA English
 FAN.CNT 1

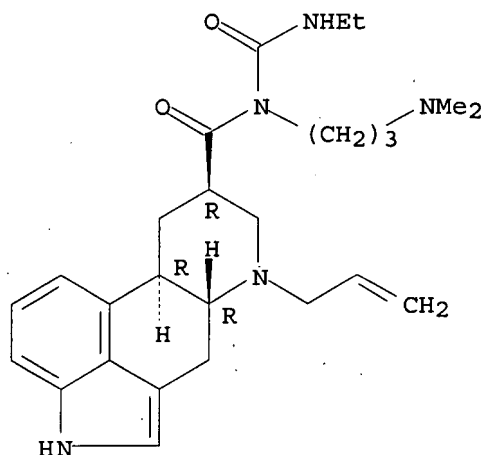
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PI	US 2004209910	A1	20041021	US 2004-827955	20040420
	US 7026483	B2	20060411		
	WO 2004094368	A2	20041104	IL 2003-155545 A	20030421
	WO 2004094368	A3	20050519	WO 2004-IB2512	20040420
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				IL 2003-155545 A	20030421
				WO 2004-IB2512 W	20040420

RE.CNT 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 6 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN
 AB A process for producing crystalline Form I of cabergoline comprises the preparation of Form V using heptane as precipitation solvent, and its exclusive conversion into crystalline Form I of cabergoline. The present crystallization process from toluene-heptane solvent system for Form V involves 'reverse addition' of toluene-cabergoline concentrate to cold heptane.
 IT 81409-90-7, Cabergoline
 RL: PEP (Physical, engineering or chemical process); PYP (Physical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
 (process for preparation of crystalline Form I of cabergoline using toluene-heptane solvent system)
 RN 81409-90-7 CAPLUS
 CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2003:757709 CAPLUS
 DN 139:281336
 TI Process for preparing crystalline Form I of cabergoline
 IN Sheikh, Ahmad Y.; Tomasi, Attilio
 PA Pharmacia Corporation, USA; Pharmacia Italia S.P.A.
 SO PCT Int. Appl., 23 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 3

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			US 2002-410163P	P	20020912
			WO 2003-EP2628	W	20030310
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			WO 2003-EP2628	W	20030310
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			US 2002-364567P	P	20020315
ZA 2004006946	A	20051005	ZA 2004-6946		20040831
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PATENT FAMILY INFORMATION:

FAN 2003:757674

PATENT NO.

KIND

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APPLICATION NO.

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				AU 2003-224665	20030310
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			US 2002-417987P	P	20021011
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US 2003232871	A1	20031218	US 2003-387173		20030312
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			NO 2004-4302	20041011
			US 2002-364567P	P 20020315
IN 2006DN07519	A	20070824	WO 2003-US7484	A 20030312
			IN 2006-DN7519	20061212
			US 2002-364567P	P 20020315
			WO 2003-US7484	W 20030312
			IN 2004-DN2444	A3 20040823

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 7 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

AB A process for producing crystalline form I of cabergoline comprises crystallization of

the desired form from a toluene/heptane or toluene/hexane mixture starting from raw cabergoline, followed by recovery and removal of the solvent from the resulting toluene solvate Form X so as to convert it into form I. The solvate form X of cabergoline and its preparation are also provided.

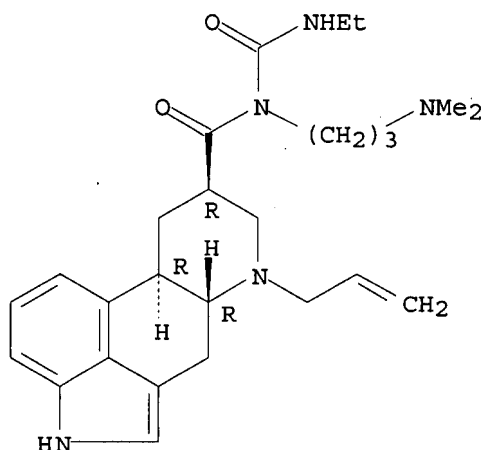
IT 81409-90-7, Cabergoline

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); PROC (Process)
(process for preparing the crystalline form I of cabergoline)

RN 81409-90-7 CAPLUS

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8 β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2003:757674 CAPLUS
DN 139:265794

TI Process for preparing the crystalline form I of cabergoline
 IN Sheikh, Ahmad Y.
 PA Pharmacia Corporation, USA
 SO PCT Int. Appl., 17 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 3

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FAN 2003:757687

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PI	WO 2003078408	A1	20030925	WO 2003-US7484	20030312
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			WO 2003-EP2628	W 20030310

L4 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

AB The crystalline form VII of cabergoline, a pharmaceutical composition containing the

form and a process for its preparation are disclosed. The process may comprise a slurry procedure by using the form I or mixture of forms I and VII in a solvent at a temperature above 30°. Thus, cabergoline was dissolved in 1,4-dioxane at 40°, the final solution was cooled to -5° and the solid was filtered and dried at 30-65°. The crystals of the form VII were obtained.

IT 81409-90-7, Cabergoline

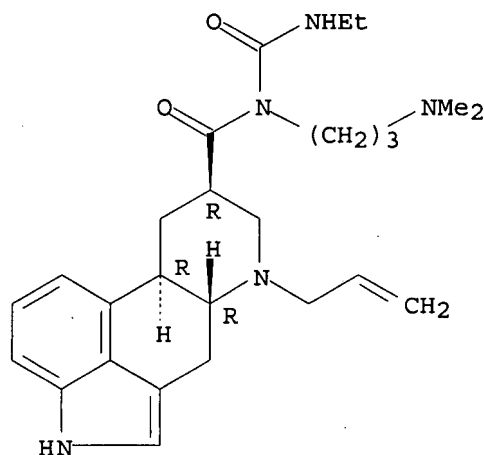
RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(crystalline form VII of cabergoline)

RN 81409-90-7 CAPLUS

CN Ergoline-8-carboxamide, N-[3-(dimethylamino)propyl]-N-[(ethylamino)carbonyl]-6-(2-propen-1-yl)-, (8β)- (CA INDEX NAME)

Absolute stereochemistry.



AN 2001:730739 CAPLUS
 DN 135:262284
 TI Crystalline form VII of cabergoline
 IN Candiani, Ilaria; Budelli, Raffaella; Pandolfi, Marco; Ungari, Mario
 PA Pharmacia & Upjohn S.p.A., Italy
 SO PCT Int. Appl., 15 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

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PI	WO 2001072746	A1	20011004	WO 2001-EP2969	20010315
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			GB 2000-7309	A 20000324
			WO 2001-EP2969	W 20010315

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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